

इंटरनेट

मानक

Disclosure to Promote the Right To Information

Whereas the Parliament of India has set out to provide a practical regime of right to information for citizens to secure access to information under the control of public authorities, in order to promote transparency and accountability in the working of every public authority, and whereas the attached publication of the Bureau of Indian Standards is of particular interest to the public, particularly disadvantaged communities and those engaged in the pursuit of education and knowledge, the attached public safety standard is made available to promote the timely dissemination of this information in an accurate manner to the public.

“जानने का अधिकार, जीने का अधिकार”

Mazdoor Kisan Shakti Sangathan

“The Right to Information, The Right to Live”

“पुराने को छोड़ नये के तरफ”

Jawaharlal Nehru

“Step Out From the Old to the New”

IS 12473-3 (1988): Chemical Analysis of Hardmetals by Flame Atomic Absorption Spectrometry, Part 3: Determination of cobalt, iron, manganese and nickel in contents from 0.01 to 0.5 percent (m/m) [MTD 25: Powder Metallurgical Materials and Products]



“ज्ञान से एक नये भारत का निर्माण”

Satyanarayan Gangaram Pitroda

“Invent a New India Using Knowledge”



“ज्ञान एक ऐसा खजाना है जो कभी चुराया नहीं जा सकता है”

Bhartrhari—Nitiśatakam

“Knowledge is such a treasure which cannot be stolen”

BLANK PAGE



Indian Standard

**CHEMICAL ANALYSIS OF
HARDMETALS BY FLAME ATOMIC
ABSORPTION SPECTROMETRY**

**PART 3 DETERMINATION OF COBALT, IRON, MANGANESE AND
NICKEL IN CONTENTS FROM 0.01 TO 0.5 PERCENT (m/m)**

UDC 669.018.25 : 543.422 [546.73]

© Copyright 1989

BUREAU OF INDIAN STANDARDS
MANAK BHAVAN, 9 BAHADUR SHAH ZAFAR MARG
NEW DELHI 110002

Indian Standard

CHEMICAL ANALYSIS OF HARDMETALS BY FLAME ATOMIC ABSORPTION SPECTROMETRY

PART 3 DETERMINATION OF COBALT, IRON, MANGANESE AND NICKEL IN CONTENTS FROM 0.01 TO 0.5 PERCENT (*m/m*)

0. FOREWORD

0.1 This Indian Standard (Part 3) was adopted by the Bureau of Indian Standards on 23 September 1988, after the draft finalized by the Powder Metallurgical Materials and Products Sectional Committee had been approved by the Structural and Metals Division Council.

0.2 Chemical analysis of hardmetals by flame atomic absorption spectrometric methods is covered in the following parts:

Part 5 Determination of cobalt, iron manganese, molybdenum, nickel, titanium and vanadium 0.5-2.0

Part 6 Determination of chromium 0.01-2.0

NOTE — The method for determination of chromium also permits determination of iron, nickel and manganese within the range of 0.01 to 2 percent.

Percent Range of Each Element (*m/m*)

Part 1 Central requirements	
Part 2 Determination of calcium, potassium, magnesium and sodium	0.001-0.2
Part 3 Determination of cobalt, iron, manganese and nickel	0.01-0.5
Part 4 Determination of molybdenum, titanium and vanadium	0.01-0.5

0.3 In the preparation of this standard, assistance has been derived from ISO 7627/3-1983 'Hardmetals — Chemical analysis by flame atomic absorption spectrometry — Part 3 Determination of cobalt, iron, manganese and nickel in contents from 0.01 to 0.5 percent (*m/m*)', issued by the International Organization for Standardization (ISO).

0.4 In reporting the result of a test or analysis made in accordance with this standard, if the final value, observed or calculated, is to be rounded off, it shall be done in accordance with IS : 2-1960*.

*Rules for rounding off numerical values (revised).

1. SCOPE

1.1 This standard (Part 3) specifies the method to be used for the determination of cobalt, iron, manganese and nickel contents in hardmetals within the range 0.01 to 0.5 percent (*m/m*) by flame atomic absorption spectrometry.

1.2 General requirements concerning the field of application, principle, interfering elements, apparatus, sampling and test report are given in Part 1 of this standard.

2. REAGENTS

2.1 During the analysis, use only reagents of analytical grade, and distilled water or water of equivalent purity.

2.1.1 Hydrofluoric Acid — $\rho = 1.12$ g/ml.

2.1.2 Concentrated Nitric Acid ($\rho = 1.42$ g/ml)

2.1.3 Ammonium Fluoride (0.1 g/ml solution)

2.1.4 Caesium Chloride (0.01 g/ml solution)

2.1.5 Standard Cobalt Solution — (1 ml = 1 mg of CO) — Weigh 1.0000 g of pure cobalt metal and dissolve in minimum quantity of dilute nitric acid (1 : 1). Add 40 ml of dilute sulphuric acid (1 : 1). Heat to white fumes. Dilute to 1 litre and mix.

Alternatively, dissolve 4.7698 g of cobalt sulphate ($\text{CoSO}_4 \cdot 7\text{H}_2\text{O}$) in water and dilute to 1 litre.

2.1.6 Standard Iron Solution (1 ml = 1 mg of Fe^{+2}) — Dissolve 7.0215 g of ammonium ferrous sulphate in 200 ml of water containing 10 ml

concentrated sulphuric acid. Dilute to 1 litre and mix.

2.1.7 Standard Manganese Solution — (1 ml = 1 mg of Mn) — Dissolve 1.0000 g of electrolytic manganese nickel (99.9 percent, *Min*) in 50 ml of dilute nitric acid (1 : 1). Cool and transfer to 1 litre volumetric flask, dilute with water up to the mark and mix.

2.1.8 Standard Nickel Solution (1 ml = 1 mg of Ni) — Dissolve 1.0000 g of nickel (99.9 percent, *Min*) in 100 ml of dilute nitric acid (1 : 1) and heat slowly; cool and transfer to 1 litre volumetric flask, dilute with water up to the mark and mix.

3. PROCEDURE

3.1 Test Portion — Weigh, to the nearest 0.001 g, approximately 1 g of the test sample. Transfer it to a 100 ml polytetrafluoroethylene beaker. Cover the beaker.

3.2 Dissolution of the Test Portion — Add 10 ml of water, 5 ml of hydrofluoric acid and then 5 ml of concentrated nitric acid, drop by drop, to the beaker containing the test portion, and heat gently until the test portion is completely dissolved. Add 10 ml each of caesium chloride and ammonium fluoride solution. Then transfer the solution to a 100 ml polyethylene volumetric flask and dilute to the mark.

3.3 Dilution Volume — Prepare the relevant dilution volume for the analysis according to Table 1.

3.3.1 Dilution Volume (100 ml = 1 g of Test Sample) — Use the solution in 3.2.

3.3.2 Dilution Volume (1 000 ml = 1 g of Test Sample) — The concentration of the solution may be reduced by a factor of 10 for instruments of higher sensitivity by transferring 10 ml of the solution in 3.2 to a 100 ml polyethylene volumetric flask. Add 10 ml of caesium chloride solution. Add 10 ml of ammonium fluoride solution and dilute to the mark.

3.4 Preparation of Calibration and Blank Solutions

3.4.1 Prepare at least six solutions according to 3.2 with a matrix composition as similar as possible to the test portion to be analysed, but without making up to volume. Then add increasing volumes of properly diluted standard solutions (2.1.5 to 2.1.8) of the elements to be determined according to the concentration ranges to be covered. Make up to 100 ml and mix.

3.4.2 Also prepare calibration solution with a diluted matrix in accordance with 3.3.2 for dilution volumes of 1 000 ml, if necessary.

3.4.3 Prepare two blank solutions (see 3.4.1) without the addition of the relevant element to be determined.

3.5 Adjustment of the Atomic Absorption Spectrometer — Ignite the flame. Optimize the response of the instrument at the wavelength given in Table 1 for the element being determined. Preheat the burner for about 5 min and then adjust the fuel and correct the burner to obtain maximum absorption while aspirating a calibration solution. Make sure that the absorbance reading is not drifting. Aspirate water and set the initial reading to zero absorbance.

TABLE 1 INSTRUMENTAL PARAMETERS AND CHARACTERISTICS OF CALIBRATION FUNCTIONS

(Clauses 3.3 and 3.5)

ELEMENT	DILUTION VOLUME (V) FOR 1 g TEST PORTION*	OXIDANT	WAVELENGTH	RECIPROCAL SENSITIVITY FOR 1 PERCENT ABSORPTION*	LINEAR RANGE*	NOTES
	ml		nm	µg/ml	Percent	
Co	100	N ₂ O	240.7	0.7	0.007 to 0.5	1
Fe	100	N ₂ O	248.3	0.3	0.003 to 0.05	
	1 000	N ₂ O	248.3	0.3	0.03 to 2.0	
Mn	100	N ₂ O	279.8	0.1	0.001 to 0.03	2
	1 000	N ₂ O	279.8	0.1	0.01 to 2.0	2
Ni	100	N ₂ O	232.0	0.3	0.003 to 0.05	3
	1 000	N ₂ O	232.0	0.3	0.05 to 5.0	3

NOTE 1 — When a broad spectral band pass is used, a single-element hollow cathode lamp is mandatory.

NOTE 2 — Use of the triplet.

NOTE 3 — Sensitivity is greatly dependent on matrix composition. Use of a narrow-spectral band pass and a single-element hollow-cathode lamp is mandatory.

*Guidelines for information only.

3.6 Atomic Absorbance Measurements

3.6.1 Aspirate first the blank solution and then the calibration and test solutions consecutively and record the readings. Aspirate water between each solution. Make at least two measurements for each solution. Solids which build up on the burner slit must be removed, otherwise they will cause a decrease of sensitivity.

3.6.2 Prepare a calibration curve by plotting the obtained absorbance values of the calibration solutions corrected for the blank against the concentrations, in mg per litre, of the element.

3.6.3 Convert the absorbance values of the test solutions corrected for the blank to mg of the element per litre by means of the calibration curve.

4. TEST RESULTS

4.1 Calculation — The element content, expressed as a percentage by mass, is given by the formula

$$\frac{C \times V}{10^4 \times m}$$

where

C = concentration, in mg per litre, of the element in the test solution;

V = dilution volume, in ml; and

m = mass, in g, of the test portion.

4.2 Permissible Tolerances — The deviation between three independent determinations shall not exceed 0.2 times the element content in percent (absolute value).

4.3 Final Result — Report the arithmetical mean of acceptable determinations rounded to the nearest value as shown in Table 2.

TABLE 2 ROUNDING OFF OF RESULTS

CONTENT	ROUND TO THE NEAREST
Percent	Percent
From 0.01 to 0.1	0.01
Over 0.1 to 0.5	0.05

4.3.1 If the element content is below 0.01 percent, report it as less than 0.01 percent.

BUREAU OF INDIAN STANDARDS

Headquarters:

Manak Bhavan, 9 Bahadur Shah Zafar Marg, NEW DELHI 110002

Telephones: 3 31 01 31, 3 31 13 75

Telegrams: Manaksanstha

(Common to all Offices)

Regional Offices:

	<i>Telephone</i>
Central : Manak Bhavan, 9 Bahadur Shah Zafar Marg, NEW DELHI 110002	{ 3 31 01 31 3 31 13 75
*Eastern : 1/14 C.I.T. Scheme VII M, V.I.P. Road, Maniktola, CALCUTTA 700054	36 24 99
Northern : SCO 445-446, Sector 35-C, CHANDIGARH 160036	{ 2 18 43 3 16 41
Southern : C.I.T. Campus, IV Cross Road, MADRAS 600113	{ 41 24 42 41 25 19 41 29 16
†Western : Manakalaya, E9 MIDC, Marol, Andheri (East), BOMBAY 400093	6 32 92 95

Branch Offices:

'Pushpak', Nurmohamed Shaikh Marg, Khanpur, AHMADABAD 380001	{ 2 63 48 2 63 49
‡Peenya Industrial Area, Ist Stage, Bangalore-Tumkur Road, BANGALORE 560058	{ 38 49 55 38 49 56
Gangotri Complex, 5th Floor, Bhadbhada Road, T. T. Nagar, BHOPAL 462003	6 67 16
Plot No. 82/83, Lewis Road, BHUBANESHWAR 751002	5 36 27
53/5 Ward No. 29, R.G. Barua Road, 5th By-lane, GUWAHATI 781003	3 31 77
5-8-56C, L.N. Gupta Marg (Nampally Station Road), HYDERABAD 500001	23 10 83
R14 Yudhister Marg, C Scheme, JAIPUR 302005	{ 6 34 71 6 98 32
117/418 B Sarvodaya Nagar, KANPUR 208005	{ 21 68 76 21 82 92
Patliputra Industrial Estate, PATNA 800013	6 23 05
T.C. No. 14/1421, University P.O., Palayam, TRIVANDRUM 695034	{ 6 21 04 6 21 17

Inspection Offices (With Sale Point):

Pushpanjali, First Floor, 205-A West High Court Road, Shankar Nagar Square, NAGPUR 440010	2 51 71
Institution of Engineers (India) Building, 1332 Shivaji Nagar, PUNE 411005	5 24 35

*Sales Office in Calcutta is at 5 Chowringhee Approach, P.O. Princep Street, Calcutta 700072 27 68 00

†Sales Office in Bombay is at Novelty Chambers, Grant Road, Bombay 400007 89 65 28

‡Sales Office in Bangalore is at Unity Building, Narasimharaja Square, Bangalore 560002 22 39 71